

K875240 TRANS/ NEEDLE GUIDEApr 15, 1988
115 days to decisionK875240 · Product code: **IYO** · Radiology
Source: <https://www.510kdatabase.net/k875240/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Imaging, Pulsed Echo, Ultrasonic (IYO)
Date received	Dec 22, 1987
Decision date	Apr 15, 1988
Days to decision	115 days
Third-party review	No

APPLICANT

Company	CIVCO Medical Instruments Co., Inc.
Location	Walker, MI, US
Contact	VICTOR WEDEL
510(k) history	29 submissions · 29 cleared · 1982-2023

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k875240/>, Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 17, 2026